

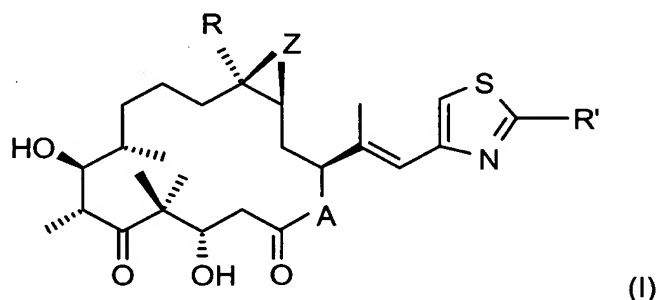
## Amendments to the Claims

### Listing of Claims:

Claims 1-19 (canceled)

Please add the following new claims:

Claim 20 (new): A combination which comprises (a) a HER-1 or a HER-2 antibody or (b) at least one antineoplastic agent selected from the group consisting of aromatase inhibitors, antiestrogens, topoisomerase I inhibitors, topoisomerase II inhibitors, microtubule active agents, protein kinase C inhibitors, anti-angiogenic compounds, gonadorelin agonists, anti-androgens, histone deacetylase inhibitors, and S-adenosylmethionine decarboxylase inhibitors and (c) an epothilone derivative of formula I



wherein A represents O or NR<sub>N</sub>, wherein R<sub>N</sub> is hydrogen or lower alkyl, R is hydrogen or lower alkyl, R' is methyl, methoxy, ethoxy, amino, methylamino, dimethylamino, aminomethyl or methylthio, and Z is O,

in which the active ingredients (a) and (b) are present in each case in free form or in the form of a pharmaceutically acceptable salt and optionally at least one pharmaceutically acceptable carrier; for simultaneous, separate or sequential use.

Claim 21 (new): Combination according to claim 20 wherein the HER-1 or HER-2 antibody is trastuzumab.

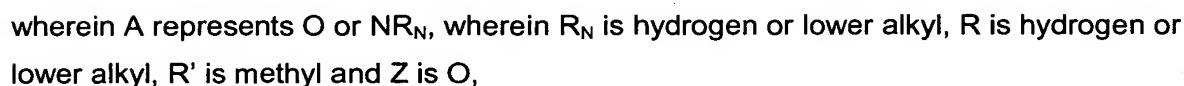
Claim 22 (new): Combination according to claim 20 wherein the antineoplastic agent is a topoisomerase I inhibitor.

Claim 24 (new): Combination according to claim 20 wherein the antineoplastic agent is an aromatase inhibitor.

Claim 26 (new): Combination according to claim 20 wherein the epothilone derivative is epothilone B.

**Claim 28 (new): A combination which comprises**

(b) an epothilone derivative of formula I

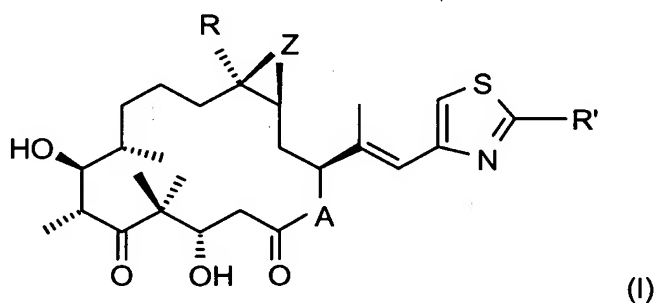


**Claim 29 (new):** Combination according to claim 28 wherein the HER-1 or HER-2 antibody is trastuzumab.

Claim 30 (new) Combination according to claim 28 wherein the epothilone derivative is epothilone B.

Claim 31 (new) Combination according to claim 28 which is a combined preparation.

Claim 32 (new): A combination which comprises (a) at least one antineoplastic agent selected from the group consisting of topoisomerase I inhibitors, topoisomerase II inhibitors, microtubule active agents, protein kinase C inhibitors, anti-angiogenic compounds, gonadorelin agonists, anti-androgens, histone deacetylase inhibitors, and S-adenosylmethionine decarboxylase inhibitors and (b) an epothilone derivative of formula I



wherein A represents O or  $\text{NR}_N$ , wherein  $\text{R}_N$  is hydrogen or lower alkyl, R is hydrogen or lower alkyl, R' is methyl, methoxy, ethoxy, amino, methylamino, dimethylamino, aminomethyl or methylthio, and Z is O,

in which the active ingredients (a) and (b) are present in each case in free form or in the form of a pharmaceutically acceptable salt and optionally at least one pharmaceutically acceptable carrier; for simultaneous, separate or sequential use.

Claim 33 (new): Combination according to claim 32 wherein the antineoplastic agent is a topoisomerase I inhibitor.

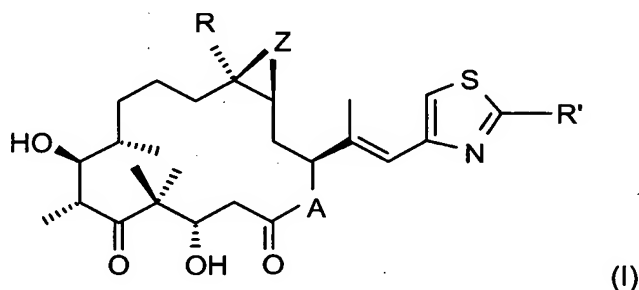
Claim 34 (new): Combination according to claim 32 wherein the antineoplastic agent is a topoisomerase II inhibitor.

Claim 35 (new): Combination according to claim 32 wherein the antineoplastic agent is a microtubule active agent.

Claim 36 (new): Combination according to claim 32 wherein the epothilone derivative is epothilone B.

Claim 37 (new): Combination according to claim 32 which is a combined preparation

Claim 38 (new): A combination which comprises (a) at least one antineoplastic agent selected from the group consisting of aromatase inhibitors and antiestrogens and (b) an epothilone derivative of formula I



wherein A represents O or  $\text{NR}_N$ , wherein  $\text{R}_N$  is hydrogen or lower alkyl, R is hydrogen or lower alkyl, R' is methyl, methoxy, ethoxy, amino, methylamino, dimethylamino, aminomethyl or methylthio, and Z is O, in which the active ingredients (a) and (b) are present in each case in free form or in the form of a pharmaceutically acceptable salt and optionally at least one pharmaceutically acceptable carrier; for simultaneous, separate or sequential use.

Claim 39 (new): Combination according to claim 38 wherein the antineoplastic agent is an aromatase inhibitor.

Claim 40 (new): Combination according to claim 38 wherein the epothilone derivative is epothilone B.

Claim 41 (new): Combination according to claim 38 which is a combined preparation.

Claim 42 (new): Combination according to claim 20 which comprises (a) a HER-1 or a HER-2 antibody or (b) at least one antineoplastic agent selected from the group consisting of aromatase inhibitors antiestrogens, topoisomerase I inhibitors, topoisomerase II inhibitors, microtubule active agents, protein kinase C inhibitors, anti-angiogenic compounds, gonadorelin agonists, anti-androgens, histone deacetylase

inhibitors, and S-adenosylmethionine decarboxylase inhibitors and (c) an epothilone derivative of formula I wherein A represents O or  $\text{NR}_N$ , wherein  $\text{R}_N$  is hydrogen or lower alkyl,  $\text{R}'$  is methyl or methylthio, R is hydrogen or lower alkyl, and Z is O, in which the active ingredients (a) and (b) are present in each case in free form or in the form of a pharmaceutically acceptable salt and optionally at least one pharmaceutically acceptable carrier; for simultaneous, separate or sequential use.

Claim 43 (new): Method of treating a warm-blooded animal having a proliferative disease comprising administering to the animal a combination according to claim 20 in a quantity which is jointly therapeutically effective against a proliferative disease and in which the compounds can also be present in the form of their pharmaceutically acceptable salts.

Claim 44 (new): A pharmaceutical composition comprising a quantity which is jointly therapeutically effective against a proliferative disease of a combination according to claim 20 and at least one pharmaceutically acceptable carrier.

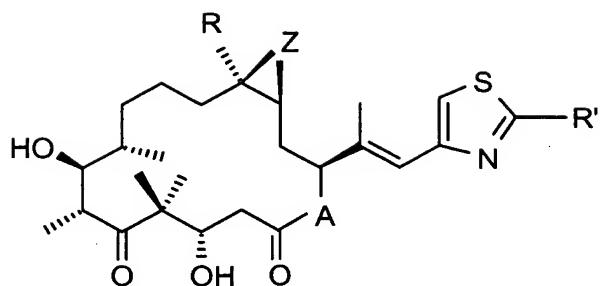
Claim 45 (new): A combination according to claim 20 for use in the treatment of a proliferative disease.

Claim 46 (new): Use of a combination according to claim 20 for the preparation of a medicament for the treatment of a proliferative disease.

Claim 47 (new): Use according to claim 45 wherein the proliferative disease is a solid tumor disease.

Claim 48 (new): Use according to claim 46 wherein the proliferative disease is a solid tumor disease.

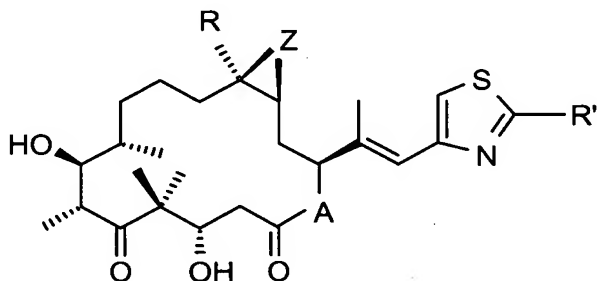
Claim 49 (new): Use of (a) a HER-1 or a HER-2 antibody or (b) at least one antineoplastic agent selected from the group consisting of aromatase inhibitors, antiestrogens, topoisomerase I inhibitors, topoisomerase II inhibitors, microtubule active agents, protein kinase C inhibitors, anti-angiogenic compounds, gonadorelin agonists, anti-androgens, histone deacetylase inhibitors, and S-adenosylmethionine decarboxylase inhibitors in combination with (c) an epothilone derivative of formula I



(I)

wherein A represents O or  $\text{NR}_N$ , wherein  $\text{R}_N$  is hydrogen or lower alkyl, R is hydrogen or lower alkyl, R' is methyl, methoxy, ethoxy, amino, methylamino, dimethylamino, aminomethyl or methylthio, and Z is O,  
for the preparation of a medicament for the treatment of a proliferative disease.

Claim 50 (new): A commercial package comprising (a) a HER-1 or a HER-2 antibody or (b) at least one antineoplastic agent selected from the group consisting of aromatase inhibitors, antiestrogens, topoisomerase I inhibitors, topoisomerase II inhibitors, microtubule active agents, protein kinase C inhibitors, anti-angiogenic compounds, gonadorelin agonists, anti-androgens, histone deacetylase inhibitors, and S-adenosylmethionine decarboxylase inhibitors and (c) an epothilone derivative of formula I



(I)

wherein A represents O or  $\text{NR}_N$ , wherein  $\text{R}_N$  is hydrogen or lower alkyl, R is hydrogen or lower alkyl, R' is methyl, methoxy, ethoxy, amino, methylamino, dimethylamino, aminomethyl or methylthio, and Z is O,  
together with instructions for simultaneous, separate or sequential use thereof in the treatment of a proliferative disease.